DETECTION OF NEURODEGENERATIVE DISEASES, *IN VITRO* SCREENING ASSAYS

BACKGROUND OF THE INVENTION

5   Diagnosis of many neurodegenerative disorders is difficult and often based solely on clinical criteria. Even when specific genetic mutations are known to cause a disease, diagnostic genetic screening may not be feasible since a variety of different point mutations in a single gene may cause the disease. The assays of the current invention permit detection of the functional outcome of these mutations by measuring altered cellular physiology of

10   blood samples taken from patients. Blood lymphocytes are transformed to lymphoblasts which are then compared to control lymphoblasts using microphysiometry. Once the lymphoblasts are available (a matter of days after receiving a blood sample), discriminative physiological testing can be performed within a few hours.

Development of therapeutics for neurodegenerative disorders is exceptionally

15   difficult. For diseases caused by known genetic mutations, transgenic animal models are being developed. However, transgenic animal models are a slow and expensive process, and their relevance and utility is variable and often limited. Even when such models are developed successfully, they do not provide an efficient means of screening potential therapeutic agents. What is needed is an *in vitro*, biologically-relevant, rapid, high-

20   throughput method for screening candidate therapeutics, both pharmacologic and genetic, in specific neurodegenerative disorders.

The primary technology applicants use to monitor cellular responses is a microphysiometer to measure a nonspecific endpoint and is broadly applicable to a variety of cellular functions. Applicants have discovered that a wide variety of different

25   pathogenic genetic defects alter cellular metabolism in a manner that can be detected using microphysiometry.

Applicants have developed a method for rapid and efficient analysis of cellular defects caused by a variety of neurodegenerative diseases, including Alzheimer's disease (AD), Huntington's disease (HD), Parkinson's disease (PD) and mitochondrial disorders,

30   such as Leber's hereditary optic neuropathy (LHON). Applicants' method employs a novel combination of cell culture and molecular biology techniques with rapid *in vitro* evaluation of cellular responses to therapeutic agents, both pharmacologic and genetic

BRIEF DESCRIPTION OF THE INVENTION

Applicants have discovered that various neurodegenerative diseases show patterns of responses in proton excretion, when the appropriate cells are incubated in the presence of inhibitors of the mitochondrial electron transport chain or in the presence of calcium ionophores, and these patterns aid in the diagnosis of diseases including Alzheimer's disease, Huntington's disease, Leber's hereditary optic neuropathy, and Parkinson's disease. These patterns also provide assays for screening compounds suitable for the treatment of a neurodegenerative disease.

Abbreviations and Definitions

AD	Alzheimer's disease
HD	Huntington's disease
LHON	Leber's hereditary optic neuropathy
PD	Parkinson's disease
TTFA	Thenoyltrifluoroacetone
MPTP	1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine
MPP^+	1-methyl-4-phenylpyridinium

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1A: A graph showing results of microphysiometry of sporadic AD lymphoblasts, which shows apparent mitochondrial defects when cells are exposed to a complex I inhibitor, rotenone.

Figure 1B: A graph showing results of microphysiometry of sporadic AD lymphoblasts, which shows apparent mitochondrial defects when cells are exposed to a complex III inhibitor, antimycin.

Figure 1C: A graph showing results of microphysiometry of sporadic AD lymphoblasts, which shows a normal response to a complex II inhibitor, malonate.

Figure 1D: A graph showing results of microphysiometry of sporadic AD lymphoblasts, which shows a normal response to a complex IV inhibitor, sodium azide.

Figure 2A: A graph showing results of microphysiometry of PC12 cells transfected with two mutant forms of PS1, both known genetic causes of AD, which shows apparent mitochondrial defects when cells are exposed to a complex I inhibitor, rotenone.

5   Figure 2B: A graph showing results of microphysiometry of PC12 cells transfected with two mutant forms of PS1, both known genetic causes of AD, which shows a normal response to a complex II inhibitor, malonate.

Figure 2C: A graph showing results of microphysiometry of PC12 cells transfected with two mutant forms of PS1, both known genetic causes of AD, which

10   shows apparent mitochondrial defects when cells are exposed to a complex III inhibitor, diuron.

Figure 2D: A graph showing results of microphysiometry of PC12 cells transfected with two mutant forms of PS1, both known genetic causes of AD, which shows a reduced response to a complex IV inhibitor, sodium azide.

15   Figure 3:   A graph showing results of microphysiometry of HD fibroblasts, which shows altered calcium homeostasis after exposure to ionomycin. This altered response to ionomycin is due to mitochondrial dysfunction since it can be reproduced in control cells by rotenone.

Figure 4A: A graph showing results of microphysiometry of LHON cells, which shows

20   that the response to a complex I inhibitor, rotenone, is virtually absent in the 3460 mutation and markedly depressed in the 14459 mutation. Inhibitor was applied as indicated by the horizontal bar.

Figure 4B: A graph showing results of microphysiometry of LHON cells, which shows a normal response to a complex II inhibitor, TTFA. Inhibitor was applied

25   as indicated by the horizontal bar.

Figure 4C: A graph showing results of microphysiometry of LHON cells, which shows that the response to a complex III inhibitor, antimycin, is also markedly depressed. Inhibitor was applied as indicated by the horizontal bar.

Figure 4D: A graph showing results of microphysiometry of LHON cells, which shows

30   that the response to IV inhibitor, sodium azide, is mildly depressed in the

3460 cells and moderately depressed in the 14459 cells. Inhibitor was applied as indicated by the horizontal bar.

Figure 4E: A graph showing results of microphysiometry of LHON cells, which shows that the response to ionomycin, a calcium ionophore, is also abnormal in both LHON cell lines. Inhibitor was applied as indicated by the horizontal bar.

Figure 5A: A graph showing results of microphysiometry of PD lymphoblasts, which shows apparent mitochondrial defects when cells are exposed to a complex I inhibitor, rotenone.

Figure 5B: A graph showing results of microphysiometry of PD lymphoblasts, which shows a normal response to a complex II inhibitor, malonate.

Figure 5C: A graph showing results of microphysiometry of PD lymphoblasts, which shows a normal response to a complex III inhibitor, antimycin.

Figure 5D: A graph showing results of microphysiometry of PD lymphoblasts, which shows a normal response to a complex IV inhibitor, sodium azide.

DETAILED DESCRIPTION OF THE INVENTION

The present invention provides a method for aiding the diagnosis of, or screening for the risk of, a neurodegenerative disease in a subject, comprising the steps of

(a) providing a quantity of lymphoblasts from a subject to be evaluated;

(b) measuring acidification rate of the lymphoblasts, in the presence or absence of one or more compounds selected from the group consisting of an inhibitor of the mitochondrial electron transport chain and a calcium ionophore; and

(c) determining the presence of a pattern of responses in the stimulation of the acidification rate to a set of said compounds, that corresponds to a neurodegenerative disease.

In one embodiment of the present invention, the acidification rate is measured with a microphysiometer.

In another embodiment of the present invention, the neurodegenerative disease is Alzheimer's disease having the pattern of responses comprising a defective response in the stimulation of the acidification rate in the presence of complex I inhibitor, a normal

response in the acidification rate in the presence of complex II inhibitor, a defective response in the stimulation of the acidification rate in the presence of complex III inhibitor, and a normal response in the acidification rate in the presence of complex IV inhibitor.

5   In another embodiment of the present invention, the neurodegenerative disease is Huntington's disease having the pattern of response comprising a defective response in acidification rate in the presence of a calcium ionophore.

In another embodiment of the present invention, the neurodegenerative disease is Leber's Hereditary Optic Neuropathy (LHON) having the pattern of responses comprising a defective response in the acidification rate in the presence of complex I inhibitor, a

10   normal response in the acidification rate in the presence of complex II inhibitor, a defective response in the acidification rate in the presence of complex III inhibitor, a defective response in the acidification rate in the presence of complex IV inhibitor, and a defective response in the acidification rate in the presence of a calcium ionophore.

In another embodiment of the present invention, the neurodegenerative disease is

15   Parkinson's disease having the pattern of responses comprising a defective response in the acidification rate in the presence of complex I inhibitor, a normal response in the presence of complex II inhibitor, a normal response in the acidification rate in the presence of complex III inhibitor, and a normal response in the acidification rate in the presence of complex IV inhibitor.

20   In another embodiment of the present invention, a compound suitable for the treatment of a neurodegenerative disease is isolated or identified by the methods of the present invention, wherein the compound substantially reverses the defective effect of one or more inhibitors of the mitochondrial electron transport chain on the acidification rate in patient cells.

25   In another embodiment of the present invention, a compound suitable for the treatment of a neurodegenerative disease is isolated or identified by the methods of the present invention, wherein the compound substantially reverses the defective effect of a calcium ionophore on the acidification rate in patient cells.

The rate at which cells excrete protons (the acidification rate) is closely linked to

30   the rate at which they convert nutrients to energy. Because energy metabolism is coupled

to ATP consumption, any event that perturbs ATP levels, such as receptor activation or calcium influx, causes a change in energy metabolism and, therefore, an alteration in proton excretion. In addition, receptor activation may also directly affect proton excretion by altering Na/H exchange. A typical microphysiometer, such as the Cytosensor[TM]

5   (Molecular Devices), uses a silicon-based sensor to measure the rate of proton excretion from 10^4 - 10^6 cells. Cells are contained in a 3 μL flow chamber in aqueous contact with the pH-sensitive surface of a light-sensitive potentiometric sensor. Flow of medium (with low buffering capacity) and test compounds are determined by a series of microprocessor-controlled pumps. The extracellular acidification rate is determined by measuring the

10   decrease in pH that occurs as acidic metabolites accumulate during brief halts in the flow of medium. The degree of stimulation of the basal acidification is calculated and plotted as a per cent of the basal acidification rate. The sensor takes a voltage measurement that is linearly related to pH. Typically, during a 30 - 60 sec halt in flow, pH will decrease less than 0.1 pH unit. Reproducible and robust responses are obtained to metabolic inhibitors,

15   growth factors and ionophores in lymphoblast lines. Because virtually any manipulation alters proton excretion, it is relatively easy to determine the types of stimuli to which disease cell lines respond abnormally.

A single microphysiometer can contain up to 8 cell chambers, thereby allowing great flexibility in experimental design. For example, a 4-point dose-response curve can be

20   generated simultaneously in control and disease cell lines. Alternatively, 8 drugs can be screened simultaneously in a disease cell line. Because the microphysiometric responses are obtained in essentially real time, this technique is suitable for relatively high-throughput screening of targeted pharmacological agents.

Alzheimer's Disease (AD)

Approximately 5-10% of AD cases are caused by autosomal genetic mutations, i.e., familial AD, and the remaining 90% of cases are of undefined cause and are termed "sporadic AD." Sporadic AD lymphoblasts have markedly blunted microphysiometric

5 responses to the mitochondrial complex I inhibitor, rotenone (Figure 1A), and to the complex III inhibitor, antimycin-A (Figure 1B). Responses to the complex II inhibitor, malonate (Figure 1C), and to the complex IV inhibitor, sodium azide (Figure 1D) do not differ substantially from controls. These results indicate that there is chronic impairment of mitochondrial function in AD and they provide a simple diagnostic test for sporadic AD.

10 In PC12 cells transfected with the most common genetic mutations that cause familial AD with onset at early age (presenilin-1), a similarly depressed response to rotenone (Figure 2A) and to the complex III inhibitor diuron (Figure 2C) were found. These results indicate that microphysiometry is suitable for the diagnosis of genetic forms of AD.

15 In addition, these results show that microphysiometry provides a method for screening potential therapeutic strategies, both pharmacologic and genetic. The ability of new drugs or genetic therapies to normalize the AD responses predicts therapeutic efficacy.

Huntington's Disease (HD)

20 A leading hypothesis of Huntington's disease pathogenesis suggests that the causative mutation in the protein, huntingtin, leads secondarily to mitochondrial dysfunction and altered calcium homeostasis. HD cells have abnormal responses to the calcium ionophore, ionomycin (Figure 3). This altered response to ionomycin is due to mitochondrial dysfunction since it can be reproduced in control cells by rotenone (Figure

25 3). These results show that microphysiometry provides a method for screening potential therapeutic strategies, both pharmacologic and genetic. The ability of new drugs or genetic therapies to normalize the HD responses predicts therapeutic efficacy.

Leber's Hereditary Optic Neuropathy (LHON)

Leber's hereditary optic neuropathy (LHON) is caused by mutations in

30 mitochondrial genes that encode subunits of complex I of the mitochondrial electron

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transport chain. Two of these mutations are the 3460 and 14459 mutations, both of which are detected by microphysiometry. The microphysiometric response to a complex I inhibitor, rotenone, is virtually absent in the 3460 mutation and markedly depressed in the 14459 mutation (Figure 4A). In contrast, there is no difference between control and

5 patient lymphoblast responses to a complex II inhibitor, TTFA (Figure 4B). Responses to a complex III inhibitor, antimycin, are also markedly depressed (Figure 4C). Responses to a complex IV inhibitor, sodium azide, are mildly depressed in the 3460 cells and moderately depressed in the 14459 cells (Figure 4D). Responses to ionomycin, a calcium ionophore, are also abnormal in both LHON cell lines (Figure 4E).

10 These results provide an alternative, non-genetic method for diagnosis of suspected LHON. These results also show that microphysiometry provides a method for screening potential therapeutic strategies, both pharmacologic and genetic. The ability of new drugs or genetic therapies to normalize the LHON responses predicts therapeutic efficacy.

Parkinson's Disease (PD)

15 Currently, the cause of PD is unknown, although one leading hypothesis proposes that altered mitochondrial complex I activity may be responsible. PD lymphoblasts have markedly blunted microphysiometric responses to the mitochondrial complex I inhibitor rotenone I (Figure 5A). Responses to the complex II inhibitor malonate (Figure 5B), to the complex III inhibitor antimycin (Figure 5C), and to the complex IV inhibitor sodium

20 azide (Figure 5D) do not differ substantially from controls. These results indicate that there is chronic impairment of mitochondial function in PD and they provide a simple diagnostic test for PD. These results also show that microphysiometry provides a method for screening potential therapeutic strategies, both pharmacologic and genetic. The ability of new drugs or genetic therapies to normalize the PD responses predicts therapeutic

25 efficacy.

Metabolic Inhibitors Suitable for the In vitro Assay of Mitochondrial Defects

There are various inhibitors of the electron transport chain in mitochondria, and they serve as useful probes of the biochemical mechanisms underlying the metabolism of mitochondria. Inhibitors of the electron transport chain inhibit the release of protons to the

30 outside of the mitochondrial cell membrane. Applicants have discovered that each

neurodegenerative disease has a distinctive response to one or more inhibitors of the electron transport chain. Applicants have selected a limited number of inhibitors, but it will be understood that many other inhibitors of mitochondrial metabolism are suitable for the *in vitro* assay of mitochondrial defects. Suitable inhibitors include, but are not limited to,

5   inhibitors of each of the complexes of the electron transport chain, e.g., complexes I, II, III or IV. Other compounds that disrupt the release of protons to the outside the mitochondrial cell membrane are also suitable for the *in vitro* assay of mitochondrial defects, e.g., calcium ionophores that disrupt cytosolic calcium levels.

Suitable inhibitors of complex I of the electron transport chain (ETC) include, but

10   are not limited to, rotenone, 1-methy-4-phenyl-1,2,3,6-tetrahydropyridine (MPTP) or 1-methyl-4-phenylpyridinium (MPP^+). One preferred complex I inhibitor is rotenone. A suitable range of concentrations of rotenone for the *in vitro* assay ranges from about 1.0 nM to about 10μM, preferably about 100 nM. A suitable range of concentrations of MPTP for the *in vitro* assay ranges from about 100 nM to about 1 mM. A suitable range

15   of concentrations of MPP^+ for the *in vitro* assay ranges from about 100 nM to about 1 mM, preferably about 10 μM.

Suitable inhibitors of complex II of the electron transport chain include, but are not limited to, thenoyltrifluoroacetone (TTFA), malonate or 3-nitropropionic acid. One preferred complex II inhibitor is malonate. A suitable range of concentrations of TTFA for

20   the *in vitro* assay varies from about 1 μM to about 100 mM. A suitable range of concentrations of malonate for the *in vitro* assay varies from about 100μM to about 100 mM, preferably about 30 mM. A suitable range of concentrations of 3-nitroproprionic acid for the *in vitro* assay varies from about 100μM to about 100 mM.

Suitable inhibitors of complex III of the electron transport chain include, but are

25   not limited to, antimycin-A or myxothiazol. One preferred complex III inhibitor is antimycin-A, also known as antimycin. A suitable range of concentrations of antimycin for the *in vitro* assay varies from about 1 μg/ml to about 1 mg/ml, preferably about 100μg/ml. A suitable range of concentrations of myxothiazol for the *in vitro* assay varies from about 100ng/ml to about 1 mg/ml.

Suitable inhibitors of complex IV of the electron transport chain include, but are not limited to, sodium azide or cyanide. One preferred inhibitor of complex IV is sodium azide. A suitable range of concentrations of sodium azide for the *in vitro* assay varies from about 100 nM to about 100 mM, preferably about 3 mM. A suitable range of

5   concentrations of cyanide for the *in vitro* assay varies from about 100 nM to about 100 mM.

Oxamic acid is preferably used to block lactate dehydrogenase. A suitable range of concentrations of oxamic acid for the *in vitro* assay varies from about 100 μM to about 100 mM, preferably about 5 mM.

10   Suitable calcium ionophores for disrupting cytosolic calcium levels include, but are not limited to, ionomycin or calcimycin (A23187). One preferred calcium ionophore is ionomycin. A suitable range of concentrations of ionomycin for the *in vitro* assay varies from about 10 nM to about 100 μM, preferably about 3μM. A suitable range of concentrations of A23187 varies from about 10 nM to about 100 μM.

15   Patient Cell Lines

Controls: Control cell lines were generated from blood samples obtained from individuals without neurologic or mitochondrial disorders.

LHON: Previous work at Emory University has identified causative mitochondrial DNA mutations in LHON patients. Well-characterized LHON patient cell lines have been

20   generated and banked. These cell lines have been characterized by molecular genetic, cellular respiration and enzymology studies. The patient cell lines were obtained from the Emory University Clinical Research Center (Emory University, Atlanta GA).

Alzheimer's disease: Lymphoblast cell lines were generated from blood samples obtained from patients followed in the Emory Alzheimer's Disease Center. All patients

25   have been evaluated by detailed neurological and neuropsychometric examinations. All individuals given a diagnosis of AD met the National Institutes of Neurological Disorders and Stroke - Alzheimer's Disease and Related Disorders Association (NINDS-ADRDA) criteria.

Huntington's disease: Cell lines (both lymphoblasts and fibroblasts) were generated

30   from blood samples obtained from patients followed in the Emory Huntington's Disease

Clinic. All patients carry a clinical diagnosis of HD based on family history, clinical symptoms and detailed neurological evaluation; diagnosis was confirmed by genetic testing.

Parkinson's disease: Cell lines were generated from blood samples obtained from patients followed in both the Emory Movement Disorders Clinic and the Alzheimer's Disease Center. The diagnosis of PD was made by Movement Disorders specialists based on clinical presentation, detailed neurologic assessment and response to anti-parkinsonian medications.

Transformation of Blood Lymphocytes to Lymphoblasts as Cell Lines for *In Vitro* Assay of Mitochondrial Defects

Patient lymphocytes are immortalized using the technique of Neitzel H, *Human Genetics* 73:320-326 (1986). In brief, the lymphocytes are removed from the buffy coat layer following Histopaque density separation. The lymphocytes are then incubated with Epstein-Barr virus (B95-8, ATCC Access Number CRL 1612) in transforming media consisting of RPMI-1640, 20% heat-inactivated fetal bovine serum (FBS), 2 μg/ml cyclosporin, 2 mM L-glutamine, and penicillin/streptomycin. Following transformation, the lymphoblasts are maintained in RPMI-1640 media supplemented with 15% heat-inactivated FBS and 110 μg/ml sodium pyruvate.

Generation of Transfected PC12 Cells as Models of AD

Rat adrenal pheochromocytoma PC12 cells were stably transfected with PS1 cDNA cloned behind the CMV promoter in pcDNA3. See, e.g., Lah, J.J. et al., *J. Neurosci.* 17:1971-1980 (1997), herein incorporated by reference. Briefly, the cells were plated onto 10 cm dishes at a density of 50,000/cm^2 and grown in a 37 °C incubator with 5% carbon dioxide. After 24 hrs, 20 μg of plasmid DNA was added to each plate in a 1 ml solution of 25 mM BES, 140 mM sodium chloride, 0.750 mM sodium phosphate, 125 mM calcium chloride, pH 6.95. The cells were then returned to a 37°C incubator with 3% carbon dioxide for an additional 18-24 hrs. The media was then replaced with fresh growth media and returned to 5% carbon dioxide. Stable PC12 transfectants were selected with 400 μg geneticin and single colonies were isolated. Clonal cell lines were

established by limiting dilution cloning to ensure growth from an individual cell, and PS1

protein expression confirmed by western blotting and immunocytochemistry.

In Vitro Assay of Mitochondrial Defects with Microphysiometry

Lymphoblasts are suspended in agarose entrapment solution at a density of $5x10^5$

5   cells / 10 µl, then are plated onto chamber inserts, and allowed to solidify for 20 minutes.

Cells are then placed into the microphysiometer recording chambers containing a low -

buffering- capacity recording buffer made of high glucose DMEM, pH 7.40, with 44.4 mM

sodium chloride replacing sodium bicarbonate in the media. In all cases, except where

stated, sodium pyruvate is present in the recording buffer at 110 µg/ml. Recording buffer

10   flows across the cells at a rate of 100 µl/min. The pump cycles on for 2 min 30 sec,

followed by a 30 sec off period. The rate of acidification is calculated as the slope of a

linear least squares fit to the relation of pH versus time during the pump's off period. Cells

are equilibrated in the running media for 50-120 min prior to drug exposure. Cells are run

in duplicate for each treatment. The pH of all final drug solutions was checked to ensure

15   that the drug did not alter the pH of the running buffer.

While the foregoing specification teaches the principles of the present invention,

with examples provided for the purpose of illustration, it will be understood that the

practice of the invention encompasses all of the usual variations, adaptations, and

modifications, as come within the scope of the following claims and its equivalents.

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